

(19)



Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 040 761 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:

04.10.2000 Bulletin 2000/40

(51) Int. Cl.⁷: **A23D 9/00, C11C 3/10,**

A23G 3/00

(21) Application number: **00200721.9**

(22) Date of filing: **29.02.2000**

(84) Designated Contracting States:

**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE**

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: **19.03.1999 EP 99200868**

(71) Applicant: **LODERS CROKLAAN B.V.**

1521 AZ Wormerveer (NL)

(72) Inventors:

- **Lantz, Ingo**
22149 Hamburg (DE)
- **Schroeder, Annette**
27749 Delmenhorst (DE)

(74) Representative:

Wurfbain, Gilles L. et al
Unilever N.V.,
Patent Division,
P.O. Box 137
3130 AC Vlaardingen (NL)

(54) **Fat mixtures**

(57) Randomized fats from triglycerides with a fatty acid composition of:

- 20-50% C18:0
- 30-60% C18:1
- 5-15% C16:0 and
- 0-15% C18:2

perform very well, when applied in food products.

EP 1 040 761 A1

Description

[0001] Natural fats are often fractionated in order to get fractions that have improved properties compared with the unfractionated fats. Examples hereof are fractions of palm oil, shea oil, coconut oil, palm kernel oil and many others. From EP 815 738 blends are known that comprise "an interesterified Shea-olein fraction". However the interesterification conditions are not given and therefore it remains unclear whether this interesterification resulted in randomisation or not. Moreover it remains unclear whether a fraction of an interesterified Sh-olein is applied or whether a fraction of Sh-olein is subjected to interesterification and the product resulting therefrom is used. The blend comprising the product is used for making a whipping cream, respectively a whipped filling fat, although in the text also the use in wrapper margarine is indicated. It is said that by using this fat post-hardening problems of the fat can be reduced. The blends are based on fats with saturated C16 and/or C18 fatty acids in it, such as palm oil or fractions thereof. A disadvantage of the upgrading of a fat by fractionation often is that in addition to the desired fraction with the required product performance also one or more fractions are obtained that are not useful for application in foods. Often these fractions are too liquid, meaning that they have too low solid fat contents if measured by an NMR pulse technique to be able to apply them successfully in foods. These fractions then have to be discarded or have to be used for less useful applications like caddle feed. It would therefore be very beneficial if for such fractions a suitable application could be found, either for the fat as such or after being converted into another fat composition. We therefore studied whether we could find ways to upgrade such (waste) oils. There are actually two different routes of improvement possible:

Route 1 consists of a hardening process which will result in a high trans fatty acid content of the improved product. Route 2 which is part of the invention consists of a randomization process which results in a product having no increase in trans fatty acids. This study resulted in the finding of novel fat compositions with beneficial properties that can be applied in many food compositions. Those novel fats were found to have higher solid fat contents than the fats they were made from. Therefore these novel fats can be used to structure eg liquid oils often used in food compositions like margarines (puff pastry or cream) or icings, while they are also very useful as frying oil either as pure fat or as a blend with one or more other fats or oils.

[0002] An example of an attempt to reuse a liquid oil is disclosed in EP 69599. According to this patent a CBE is made from shea oil by fractionation. The olein fraction formed as side product is up-graded by subjecting it to a directed 1,3 enzymic interesterification. By this process the fatty acid groups bonded to the 1,3 positions of the triglycerides are redistributed and some more of the SOS compound is formed. This is isolated by fractionation. Still an olein fraction is obtained as side product for which no use exists.

[0003] Thus our invention concerns in the first instance a fat composition having a triglyceride composition, corresponding with a randomized, interesterified fat having a fatty acid composition of:

20-50 wt% C18:0
30-60 wt% C18:1
5-15 wt% C16:0 and
0-15 wt% C18:2

[0004] Our process leads to a different triglyceride composition than obtained by the process according to EP 69599, because in that process only the fatty acids in the 1 and 3 positions are interexchanged, whereas in our process all fatty acid groups present in the fat participate in the interesterification.

[0005] Preferred fat compositions are fat compositions, wherein the composition comprises

4 to 10 wt% of SSS
5 to 15 wt% of SUS
10 to 20 wt% of SSU
15 to 30 wt% of SUU
8 to 20 wt% of USU
12 to 30 wt% of UUU-triglycerides

preferably 4-8 wt% S3, 18-32 wt% S2U, 35-48 wt% SU2, 14-20 wt% U3
(S=saturated fatty acid with 16-18 C-atoms and U=unsaturated fatty acid with 18 C-atoms)
and which preferred fat compositions display a fatty acid composition (by FAME) of:

20-50 wt% C18:0, preferably 25-35 wt%
30-60 wt% C18:1, preferably 45-56 wt%

5-15 wt% C16:0, preferably 6-10 wt%

0-15 wt% C18:2, preferably 5-10 wt%, while the composition displays a solid fat content at the temperature indicated of:

5 N10=10-30
N20= 8-20
N30= 5-8
N35 < 5.

10 [0006] A particularly preferred fat is a fat that is obtained as a randomised, interesterified Shea oil olein. Shea oil olein is obtained as a side stream during the production of Shea oil stearin (a well known cocoa butter equivalent) from Shea oil. However so far no useful application of Shea oil olein was known, without an increase of trans fatty acids.

[0007] The fat according to the invention can be applied as such, however it is also possible to use blends of this fat with other fats or oils. In that instance the novel fats can act as structuring fat for the other fats. Therefore the invention also concerns blends of triglycerides, comprising 10 to 90 wt% of a fat composition according to claims 1-3 and 10 to 90 wt% of another fat, preferably selected from the group consisting of liquid oils (sunflower oil, rape seed oil, soybean oil, arachidic oil), tropical oils (coconut oil, palm kernel oil, palm oil), fractions thereof, or fully or partially hydrogenated products or interesterified mixes of them. In particular the use of the hard or fully hardened vegetable fat (fractions) leads to good results.

20 [0008] Preferred examples of other fats are fats that are liquid at ambient temperature (i.e having an N20 < 5, such as sunflower oil, soybean oil, olive oil etc.) and/or having relatively high contents of C12/C14 fatty acid residues (such as coconut oil; or palm kernel fat) and/or having substantial amounts of C22 or higher fatty acid residues in it (such as arachidic oils or rape oils).

[0009] Another main advantage of our novel fats is that they are derived from natural fats and that thus the products obtained after interesterification are about free of trans fatty acids. Trans fatty acids are considered less healthy nowadays and thus our novel fats can be considered as healthier than the fats with a relatively high trans content. Our invention therefore is also directed to fat compositions, wherein the composition has a trans fatty acid content of less than 10 wt%, preferably less than 5 wt%, most preferably less than 2 wt%. Trans content being defined here as the total of the trans fatty acids over the total number of unsaturated bonds in the unsaturated fatty acids present.

30 [0010] Very beneficial applications of our novel fats or of blends containing them are the use of these fats or blends in icings, resulting in increased plasticity and decreased trans fatty acids compared to common icings. Icing compositions comprising

30-70 wt% of a fat

0-30 wt% of powder products or mass (i.e. SMP, fruit, cacao etc.)

10-60 wt% of one or more sugars or sugar alcohols or mixtures of them

0.1-3 wt% of an emulsifier, wherein the fat comprises 10-100 % of the fat according to claims 1-3.

40 [0011] Our fats can also be used successfully in margarine compositions, in particular in puff pastry margarines and in cream margarines. By using our fats or our blends puff pastry margarines can be made comprising a fat emulsion with a fat content of 40-90 wt%, wherein the fat is a fat or a fat blend according to claims 1-5, while the total emulsion displays a Stevens hardness C at 20 °C of 1000 to 4000 g., or cream margarines can be made comprising a fat emulsion with a fat content of 40-90 wt% wherein the fat is a fat or a fat blend according to claims 1-5 while the emulsion displays a Stevens hardness C at 20 °C of 400 to 1500 g. Stevens hardness being measured using a Stevens Texture

45 analyzer with a probe of 12 mm and using a penetration speed of 1 mm/s and a penetration depth of 10 mm.
[0012] The fats or fat blends according to the invention can also be used for the preparation of shortenings or frying fats. Hereto some silicon additive and/or emulsifier can be incorporated into the fat or fat blend. Any known silicon additive or emulsifier can be applied, although we prefer to use types of siloxanes and/or all kinds of lecithines and/or monoglyceride mixes. Therefore in another embodiment our invention also concerns with a shortening or a frying fat composition comprising 10 to 100 wt% of fat and at least one of: (i) silicon additives and (ii) emulsifier such that the total of (i) + (ii) > 0% and flavours in amounts of 0-1 wt% of silicon additives and/or 0-2 wt% of emulsifier and/or 0-2 wt% of flavour, wherein the fat is a fat or fat blend according to claims 1-5. Our novel fats can be obtained by the following process

55 - a natural vegetable fat or a blend of natural fats is selected with a FAME of:

20-50 wt% of C18:0

30-60 wt% of C18:1

5-15 wt% of C16:0 and
0-15 wt% of C18:2

- which fat is subjected to a randomisation using a base as catalyst
- 5 - whereupon a randomised fat is isolated from the reaction mix after washing.

According to a last embodiment our invention concerns also the use of a fat composition with the composition according to claims 1-5, wherein the composition is applied as a frying fat.

[0013] It should be noted that the N-values indicated in this application were measured on fats that were subjected to the following T-regime:

melt fat at 80 °C; adjust temperature to 60 °C; cool fat until 0 °C and keep fat at 0 °C for 60 min, heat fat to measurement temperature and keep it at this temperature for 30 min and measure N-value at the temperature indicated.

15 EXAMPLES

Example 1: Preparation of In SH-f from Shf:

[0014] Shea olein was neutralized and dried before interesterification (FFA after treatment less than 0,05 %, moisture less than 0,07%). Interesterification was divided into different steps:

- heating of the raw material up to 110 °C under vacuum condition
- adding 0,2 % of sodiummethyle (catalyst). Temperature was kept constant for 15 minutes under stirring conditions. During this time interesterification took place.
- 25 - cooling to 95 °C
- soap deposit by adding water
- drying under vacuum
- bleaching and deodorisation under standard conditions

30 [0015] Overall fame analysis on triglycerids of randomized Shea olein was carried out on triglyceride recovered from an alumina column, following the standard operating procedures.

FAME	OVERALL
C 12:0	0,4
C 14:0	0,2
C 16:0	7,8
C 16:1	0,1
C 18:0	28,1
C 18:1	53,2
C 18:2	7,9
C 18:3	0,2
C 20:0	1,1
C 20:1	0,5
C 22:0	0,1

Solids content of SH-f and inSHf:

[0016]

SOLIDS:	N10	N20	N30	N35
SHf	1,2	0,7	0,5	0,1
inSHf	15,9	9,6	15,4	3,5

Triglyceride distribution	SHf	inSHf
SSS	1,5	5,6
SOS	19,5	7,8
SSO	2,6	15,7
SLnS	4,7	1,1
SSLn	0,2	2,3
OSO	1,2	11
SOO	34,6	22
SOLn	11,7	9,7
OOO	15,3	15,5
Rest	8,7	9,3

Example 2: Preparation of blends comprising inSHf:

[0017] Fully refined and randomized sheaolein was blended with fully refined other fat components in the following ratio (wt %) :

Blend	palm oil	palm-stearine mp 52	partly hardened palm-oil, mp 42	LE-rape seed oil	partly hardened LE-rape seed oil, mp 30	rand-omized shea olein	coconut oil	palm-ker-nel oil
1		5	45	10		40		
2		60		15		25		
3	30	8	36	13		13		
4					100			
5							100	
6					90	10		
7						10		90
8						10	90	
9	60	10				30		

(continued)

Blend	palm oil	palm-stearine mp 52	partly hardened palm-oil, mp 42	LE-rape seed oil	partly hardened LE-rape seed oil, mp 30	rand-omized shea olein	coconut oil	palm-ker-nel oil
10	100							

[0018] These blends were used for the applications, mentioned in examples 3-6.

Example 3: Application of fat-blends in puff pastry

3.1: Preparation of puff pastry fat

[0019]

FAME	Blend 1	Blend 2
C 12:0	1,4	0,9
C 14:0	1,0	1,1
C 16:0	26,0	38,9
C 16:1	0,2	0,2
C 18:0	16,8	8,9
C 18:1	45,4	38,1
C 18:2	6,1	8,5
C 18:3	1,2	1,5
C 20:0	0,6	0,5
C 20:1	0,4	0,3
C 22:0	0,1	0,1
Trans-Fatty Acid (%)	6,6	virtually free
Solid content 20°C (N20)	38,0	39,0
Solid content 30°C (N30)	20,8	23,8
Solid content 35°C (N35)	13,9	17,4
Iodine Value	71,1	65,6

[0020] The N-values were measured by means of NMR (one hour stabilisation).

[0021] From the fat blends 1 and 2 a puff pastry margarine was manufactured. The margarine had the following composition (wt % on total composition):

Fat phase: 82 %

81,5 % fat

0,2 % Lecithine Adlec

0,3 % Mono-Di-glyceride Admul4203

trace beta-Carotene and flavour

Water phase: 18 %

17,4 water
 0,4 % whey powder
 0,2 % citric acid

5 [0022] The margarine was processed at laboratory scale through a conventional A-A-B sequence with a throughput of 95 l/hr., an exit temperature on the first A-unit (260 rpm) of 21 °C, an exit temperature on the second A-unit (270 rpm) of 18 °C and a packaging temperature of 24 °C. Line pressure was 30 bar. The margarine was stored at 20 °C.

[0023] Unexpected good puff pastry fats resulted, which had good plasticity to make them suitable for a lamination fat. The products exhibited the following „Stevens“ values after 1 week (12 mm probe, texture analyzer): Fat with blend
 10 1: 3213g, fat with blend 2: 2845 g.

3.2. Preparation of puff pastries:

[0024] The resulting puff pastry margarine were used for preparing pastries according to the following method:

15 [0025] Dough composition (total weight in grams):

20	Flour	1000
	Salt	15
	Sugar	30
	Water	600
25	Puff Pastry Fat	1000

[0026] Dough was kneaded for 3,5 minutes and rested for 10 minutes before lamination. Dough temperature 20 °C. Lamination was done in a two step procedure where after the first lamination a 10 minutes resting time was used and
 30 after the second lamination a resting time of 1 hour. From this laminated dough pastries of 35 grams were formed and baked in a conventional baking oven (Debag) for 28 minutes at 190 °C (75 % rel.moisture).

[0027] The products exhibited beneficial and advantageous results in:

- general impression
- 35 - turn-over
- patty height (Blend 1: 3,9 cm, Blend 2: 4,1 cm)

Example 4: Application of a fat blend in baking creams

40 4.1: Preparation of a cream baking fat

[0028]

45	FAME	Blend 3
	C 12:0	0,6
	C 14:0	0,9
50	C 16:0	35,4
	C 16:1	0,4
	C 18:0	9,8
55	C 18:1	43,1
	C 18:2	7,1
	C 18:3	1,1

(continued)

FAME	Blend 3
C 20:0	0,4
C 20:1	0,3
C 22:0	0,1
Trans-Fatty Acid (%)	5,8
Solid content 20°C (N20)	36,7
Solid content 30°C (N30)	18,9
Solid content 35°C (N35)	11,8
Iodine Value	51,8

[0029] The N-values were measured by means of NMR (one hour stabilisation).

[0030] From this fat blend a baking cream was manufactured. The cream had the following composition (wt% on total composition):

40,0 % Fat
 35,3 % Sugar
 20,0 % Water
 4,0 % Mono-Di-glyceride Admul 4204
 0,5 % Salt
 0,2 % Tartaric acid
 traces of colouring agents and flavours

[0031] The baking cream was processed at laboratory scale through a conventional A-A-B sequence with a throughput of 85 l/hr., an exit temperature on the first A-unit (230 rpm) of 22 °C, an exit temperature on the second A-unit (270 rpm) of 20 °C and a packaging temperature of 23 °C. Line pressure was 30 bar. The baking cream was stored at 20 °C.

[0032] The baking cream obtained had good properties to make them suitable for baking application. The product exhibited a „Stevens“ values after 1 week (12 mm probe, texture analyzer) of 1638 g.

4.2.: Preparation of Pan-baked white bread:

[0033] The baking cream was also used for preparing a Pan-baked white bread according to the following method:

[0034] Dough composition (total weight in grams):

Flour	3000
Salt	45
Yeast	240
Water	1500
Cream	720

[0035] Dough was kneaded in a spiral mixer for 8 minutes (1 min. slow, 7 min. fast) and rested for 30 minutes (two times a rough folding after 15 minutes each). Dough temperature was 26,1 °C. Prooving was done for 5 minutes at 36 °C before folding procedure and for another 45 minutes after the folding procedure. Folding procedure was only used to get the right shape of the dough by using a lamination machine. Pan-baked white bread was baked in a conventional baking oven for 35 minutes at 210 °C (75 % rel. moisture).

[0036] The baked product had an excellent general impression, especially the good volume (1900 g/l) and the

improved elasticity of the crumb is advantageous for this kind of products.

Example 5: Application of fat-blends in icings / glazers / coatings

5.1: Preparation of icings / glazers / coatings

[0037]

FAME	Blend 4	Blend 5	Blend 6	Blend 7	Blend 8
C 12:0	0,1	45,9	0,1	47,6	41,3
C 14:0	0,1	17,2	0,1	16,0	15,5
C 16:0	5,1	9,11	5,3	8,2	8,1
C 16:1	0,1	0,1	0,1	0,1	0,1
C 18:0	6,0	2,7	8,2	5,0	5,2
C 18:1	77,2	7,6	74,8	18,8	12,2
C 18:2	5,9	2,2	6,1	2,7	2,8
C 18:3	0,3	0,1	0,3	0,1	0,1
C 20:0	0,7	0,1	0,8	0,2	0,2
C 20:1	0,2	0,1	0,3	0,1	0,1
C 22:0	0,4	0,1	0,4	0,1	0,1
Trans-Fatty Acid (%)	57	virtually free	51,4	virtually free	virtually free
Solid content 20 °C (N20)	44,1	34,2	40,0	33,6	23,1
Solid content 30 °C (N30)	5,2	0,3	5,8	0,2	0,4
Solid content 35 °C (N35)	0,1	0,1	1,0	0	0

[0038] The N-values were measured by means of NMR (one hour stabilisation).

[0039] From these fat blends icings were manufactured with following composition (wt%):

57 % icing sugar
40 % fat
2,5 % skim milk powder
0,5 % lecithine Bolec Z
traces of colouring agents and flavours

[0040] The icings were processed on a Fryma mill (angular space 0,3 mm) at 45 °C. After crystallization at 10 °C for 24 hours, the samples were stored at 20 °C.

5.2: Assessment of icings / glazers / coatings

[0041] The icings were analyzed concerning hardness (force) and flexibility (distance) on a texture analyzer (Stable Micro Systems). For preparation the icing was melted at 40 °C, formed into a small bar (volume 10 cm³) and stored at minus 20 °C.

[0042] Analysis of the physical properties of icings (average value of 5 measurements / 5 bars):

	Blend 4	Blend 5	Blend 6	Blend 7	Blend 8
Force (g)	2500	2500	2400	3500	3700
Distance (mm)	0,55	0,65	0,7	0,55	0,45

[0043] The results show that the product properties concerning flexibility and hardness can be modified by using randomized shea olein (compare blends 4 and 6; respectively blends 5 and 8). Especially the application of the randomized shea olein in a blend for icings in the ice-cream industry is advantageous because of the good flexibility at minus 20 °C.

Example 6: Application of fat-blends as frying fat

6.1: Preparation of frying fat

[0044]

FAME	Blend 9	Blend 10
C 12:0	1,1	0,2
C 14:0	1,0	1,0
C 16:0	36,8	44,2
C 16:1	0,2	0,2
C 18:0	8,3	4,3
C 18:1	42,6	39,1
C 18:2	8,5	9,9
C 18:3	0,3	0,2
C 20:0	0,5	0,4
C 20:1	0,2	0,1
C 22:0	0,1	0,1
Trans-Fatty Acid (%)	virtually free	virtually free

[0045] The components of the fat blends were mixed and used for frying analysis

6.2.: Assessment of frying fats

[0046] The fats (each 13,5 kg) were filled in a frying pan and every 2 hours 1 kg of raw French fries were added and fried at 180 °C. This was done 4 times a day for 6 days in total. Samples of the fats were analysed in laboratory.

	Blend 9	Blend 10
Start (0 hr): Free Fatty Acid (%)	0,05	0,04
Start (0 hr): Colour Lovibond Red	2,5	2,3

(continued)

	Blend 9	Blend 10
Start (0 hr): Colour Lovibond Yellow	26,0	25,0
Start (0 hr): Smoke point (°C)	224	228
Start (0 hr): apolar parts (%)	90,7	91,8
24 hr: Free Fatty Acid (%)	0,29	0,31
24 hr: Colour Lovibond Red	4,3	2,7
24 hr: Colour Lovibond Yellow	58,0	35,0
24 hr: Smoke point (°C)	192	172
24 hr: apolar parts (%)	78,5	74,7
48 hr: Free Fatty Acid (%)	0,70	0,72
48 hr: Colour Lovibond Red	n.d.	6,2
48 hr: Colour Lovibond Yellow	n.d.	> 70
48 hr: Smoke point (°C)	188	168
48 hr: apolar parts (%)	67,5	64,0

[0047] Results show that by using randomized shea olein (=blend the frying properties can be improved. Especially the chemical decomposition and polymerisation of the fat components is lowered compared to a standard frying fat (blend 10) resulting in a better smoke point.

[0048] Although from a chemical point of view the use of randomized shea olein is advantageous, the change in colour is different compared to the standard frying fat. Nevertheless the fried product (French fries) did not have any off-taste or colouring defect.

Claims

1. Fat composition having a triglyceride composition, corresponding with a randomized, interesterified fat having a fatty acid composition of:

20-50 wt% C18 : 0
 30-60 wt% C18 : 1
 5-15 wt% C16 : 0 and
 0-15 wt% C18 : 2

2. Fat composition according to claim 1, wherein the composition comprises

4 to 10 wt% of SSS
 5 to 15 wt% of SUS
 10 to 20 wt% of SSU
 15 to 30 wt% of SUU
 8 to 20 wt% of USU
 12 to 30 wt% of UUU-triglycerides

preferably 4-8 wt% S3, 18-32 wt% S2U, 35-48 wt% SU2, 14-20 wt% U3

S=saturated fatty acid with 16-18 C-atoms and U=unsaturated fatty acid with 18 C-atoms and displaying a fatty acid composition (by FAME) of:

20-50 wt% C18:0, preferably 25-35 wt%
 30-60 wt% C18:1, preferably 45-56 wt%
 5-15 wt% C16:0, preferably 6-10 wt%
 0-15 wt% C18:2, preferably 5-10 wt%, while the composition displays a solid fat content at the temperature indicated of:

N10=10-30

N20= 8-20

N30= 5-8

N35 < 5

5

3. Fat composition according to claims 1-2, wherein the composition is randomized, interesterified Shea olein.

10

4. Blend of triglycerides comprising 10 to 90 wt% of a fat composition according to claims 1-3 and 10 to 90 wt% of another fat, preferably selected from the group consisting of liquid oils (sunflower oil, rape seed oil, soybean oil, arachidic oil), tropical oils (coconut oil, palm kernel oil, palm oil), fractions thereof, or fully or partially hydrogenated products or interesterified mixes of them.

15

5. Fat composition according to claims 1-4, wherein the composition has a trans fatty acid content of less than 10 wt%, preferably less than 5 wt%, most preferably less than 2 wt%.

6. Icing composition comprising:

20

30-70 wt% of a fat

0-30 wt% of powder products or mass (i.e. SMP, fruit, cacao etc.)

10-60 wt% of one or more sugars or sugar alcohols or mixtures of them

0.1-3 wt% of an emulsifier, wherein the fat comprises 10-100 % of the fat according to claims 1-3.

25

7. Puff pastry margarine comprising a fat emulsion with a fat content of 40-90 wt%, wherein the fat is a fat or a fat blend according to claims 1-5, while the total emulsion displays a Stevens hardness C at 20 °C of 1000 to 4000 g.

8. Cream margarine, comprising a fat emulsion with a fat content of 40 -90 wt% wherein the fat is a fat or a fat blend according to claims 1-5 while the emulsion displays a Stevens hardness C at 20 °C of 400 to 1500 g.

30

9. Shortening and frying fat composition comprising 10 to 100 wt% of fat and at least one of silicon additives and emulsifier in amounts of 0-1 wt% of silicon additives and/or 0-2 wt% of emulsifier, wherein the fat is a fat or fat blend according to claims 1-5.

10. Process for the preparation of a fat with the composition according to claims 1-5 wherein;

35

- a natural vegetable fat or a blend of natural fats is selected with a FAME of:

20-50 wt% of C18:0

30-60 wt% of C18:1

5-15 wt% of C16:0 and

40

0-15 wt% of C18:2

- which fat is subjected to a randomisation using a base as catalyst

- whereupon a randomised fat is isolated from the reaction mix after washing.

45

11. Use of a fat composition with the composition according to claims 1-5, wherein the composition is applied as a frying fat.

50

55



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 20 0721

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	US 4 214 012 A (AINGER ET AL.) 22 July 1980 (1980-07-22)	1	A23D9/00 C11C3/10
A	* example 8; table IV *	2-5	A23G3/00
X	"Bailey's industrial oil & fat products, fifth edition, vol. 3 - Edible oil and fat products: products and application technology" 1996, JOHN WILEY & SONS, INC., NEW YORK, NY, US XP002112226 * page 382; table 9.12 * * column SAL *	1	
A	EP 0 519 542 A (UNILEVER) 23 December 1992 (1992-12-23) * claims; examples *	1-5	
D, A	EP 0 069 599 A (UNILEVER) 12 January 1983 (1983-01-12) * example 1 *	1-5	
D, A	EP 0 815 738 A (UNILEVER) 7 January 1998 (1998-01-07) * claims *	1-11	TECHNICAL FIELDS SEARCHED (Int.Cl.7) A23D C11C A23G
A	EP 0 245 076 A (UNILEVER) 11 November 1987 (1987-11-11) * the whole document *	1, 2	
A	US 5 190 868 A (KOKUSHO ET AL.) 2 March 1993 (1993-03-02) * example 15 *	1, 3	
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 30 June 2000	Examiner Lepretre, F
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

EPO FORM 1503 03.82 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 20 0721

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

30-06-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4214012 A	22-07-1980	GB 1382573 A	05-02-1975
		AT 337520 B	11-07-1977
		AT 7872 A	15-10-1976
		AU 473397 B	24-06-1976
		AU 3756372 A	05-07-1973
		BE 777783 A	02-05-1972
		CA 956503 A	22-10-1974
		CH 567369 A	15-10-1975
		DE 2200461 A	10-08-1972
		ES 398640 A	16-06-1975
		FR 2121601 A	25-08-1972
		IE 35950 B	07-07-1976
		IT 1048393 B	20-11-1980
		NL 7200156 A,B,	10-07-1972
		SE 381554 B	15-12-1975
		US 4292338 A	29-09-1981
		ZA 7200057 A	29-08-1973
EP 519542 A	23-12-1992	NONE	
EP 69599 A	12-01-1983	AT 22113 T	15-09-1986
		AU 553171 B	03-07-1986
		AU 8682282 A	02-02-1983
		CA 1241227 A	30-08-1988
		DE 3273180 D	16-10-1986
		DK 112383 A,B,	07-03-1983
		ES 513828 D	16-08-1983
		ES 8308356 A	16-11-1983
		WO 8300161 A	20-01-1983
		IE 53399 B	09-11-1988
		JP 1039757 B	23-08-1989
		JP 58501066 T	07-07-1983
		MY 35387 A	31-12-1987
EP 815738 A	07-01-1998	SG 21187 G	19-02-1988
		ZA 8204872 A	29-02-1984
EP 815738 A	07-01-1998	CA 2207358 A	26-12-1997
		JP 10056964 A	03-03-1998
		US 5935627 A	10-08-1999
EP 245076 A	11-11-1987		
		GB 2190394 A	18-11-1987
		AU 596411 B	03-05-1990
		AU 7248187 A	12-11-1987
		JP 4030836 B	22-05-1992
		JP 63017697 A	25-01-1988
		SE 467860 B	28-09-1992

EPO FORM P469

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 20 0721

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

30-06-2000

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 245076 A		SE 8701847 A	07-11-1987
		ZA 8703239 A	25-01-1989
US 5190868 A	02-03-1993	JP 1137988 A	30-05-1989
		JP 2719667 B	25-02-1998
		DE 3853656 D	01-06-1995
		DE 3853656 T	18-01-1996
		EP 0305901 A	08-03-1989